This document is scheduled to be published in the Federal Register on 03/15/2016 and available online at <a href="http://federalregister.gov/a/2016-05796">http://federalregister.gov/a/2016-05796</a>, and on <a href="http://federalregister.gov/a/2016-05796">FDsys.gov</a>

## **DEPARTMENT OF JUSTICE**

[OMB Number 1117-0008]

Agency Information Collection Activities; Proposed eCollection, eComments Requested;

Extension Without Change of a Previously Approved Collection, Application for Procurement

Quota for a Controlled Substance and for Ephedrine, Pseudoephedrine, and

Phenylpropanolamine, DEA Form 250

**AGENCY:** Drug Enforcement Administration, Department of Justice.

**ACTION:** 30-Day Notice.

**SUMMARY**: The Department of Justice (DOJ), Drug Enforcement Administration (DEA), will be submitting the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. This proposed information collection was previously published in the Federal Register at 81 FR 1219, on January 11, 2016, allowing for a 60 day comment period.

<u>DATES</u>: Comments are encouraged and will be accepted for an additional 30 days until [INSERT DATE 30 DAYS AFTER DATE OF PUBLICATION IN THE FEDERAL REGISTER].

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**FOR FURTHER INFORMATION CONTACT**: If you have comments on the estimated public burden or associated response time, suggestions, or need a copy of the proposed information collection instrument with instructions or additional information, please contact Barbara J. Boockholdt, Office of Diversion Control, Drug Enforcement Administration; Mailing Address: 8701 Morrissette Drive, Springfield, Virginia 22152; Telephone: (202) 598-6812. Written comments and/or suggestions can also be directed to the Office of Management and Budget, Office of Information and Regulatory Affairs, Attention Department of Justice Desk Officer, Washington, DC 20503 or sent to OIRA\_submissions@omb.eop.gov.

**SUPPLEMENTARY INFORMATION**: Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address one or more of the following four points:

- Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;
- Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;
- Evaluate whether and if so how the quality, utility, and clarity of the information proposed to be collected can be enhanced; and

Minimize the burden of the collection of information on those who are to respond,
 including through the use of appropriate automated, electronic, mechanical, or other
 forms of information technology, e.g., permitting electronic submission of responses.

## Overview of this information collection:

- 1. Type of Information Collection: Extension of a currently approved collection.
- Title of the Form/Collection: Application for Procurement Quota for Controlled Substance and for Ephedrine, Pseudoephedrine, and Phenylpropanolamine (DEA Form 250).
- 3. The agency form number, if any, and the applicable component of the Department sponsoring the collection: Form Number: DEA Form 250. The applicable component within the Department of Justice is the Drug Enforcement Administration, Office of Diversion Control.
- 4. Affected public who will be asked or required to respond, as well as a brief abstract:

  Affected public (Primary): Business or other for-profit.

Affected public (Other): None.

Abstract: Any United States companies that desire to use any basic class of controlled substances listed in schedule I or II or the List I chemicals ephedrine, pseudoephedrine, or phenylpropanolamine for purposes of manufacturing during the

next calendar year shall apply on DEA Form 250 for a procurement quota for such

class.

5. An estimate of the total number of respondents and the amount of time estimated for

an average respondent to respond: The DEA estimates that each form takes 0.5

hours to complete. In total, 417 respondents submit 2,960 responses, with each

response taking 0.5 hours to complete.

6. An estimate of the total public burden (in hours) associated with the proposed

collection: The DEA estimates that this collection takes 1,480 annual burden hours.

If additional information is required please contact: Jerri Murray, Department Clearance Officer,

United States Department of Justice, Justice Management Division, Policy and Planning Staff,

Two Constitution Square, 145 N Street NE, Suite 3E.405B, Washington, DC 20530.

Dated: March 10, 2016.

Jerri Murray,

Department Clearance Officer for PRA,

U.S. Department of Justice.

Billing Code: 4410-09-P

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